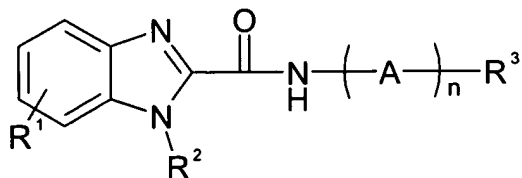


1. (currently amended) A method of treating a mammal having precancerous lesions comprising administering a pharmacologically effective amount of a compound of the following formula or pharmaceutically acceptable salt thereof:



wherein [ $R_1$  is]  $R^1$  is a hydrogen atom or a halogen atom;

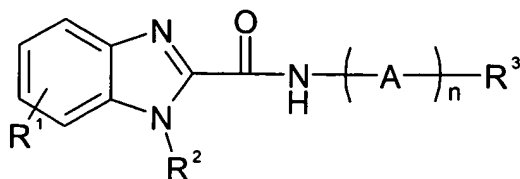
[ $R_2$  is]  $R^2$  is a phenyl-lower alkyl group;

[ $R_3$  is]  $R^3$  is a heterocyclic group selected from the group consisting of an indolyl group, indolynyl group, 1H-indazolyl group, 2(1H)-quinolinonyl group, 3,4-dihydro-2(1H)-quinolinonyl group and 3,4-dihydro-1,4(2H)-benzoxazinyl group, said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a group of the formula  $-B-R^4$ , (wherein B is a lower alkylene group;  $R^4$  is a 5-to 11-membered saturated or unsaturated heterocyclic group of single ring or binary ring, having 1 to 4 hetero atoms selected from the group consisting of a nitrogen atom, oxygen atom and sulfur atom, (said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a halogen atom, a lower alkyl group, a lower alkoxy group and oxo group) or a group of the formula  $-NR^5R^6$  (wherein  $R^5$  and  $R^6$  are each the same or different, and a hydrogen atom, a lower alkyl group, a cycloalkyl group, a pyridyl-carbonyl group, an isoxazolylcarbonyl group which may have 1 to 3 lower alkyl groups as the substituents, a pyrrolylcarbonyl group or an amino-substituted lower alkyl group which may have a lower alkyl group as the substituent; further  $R^5$  and  $R^6$  may form 5- to 6-membered saturated heterocyclic group by combining to each other, together with the adjacent nitrogen atom being bonded thereto, further with or without other nitrogen atom or oxygen atom; said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a hydroxy group and a phenyl group); a lower alkenyl group; a lower alkoxy carbonyl group; a phenoxy-lower alkyl group which may have cyano group as the substituents; a halogen-substituted lower alkyl group; and a lower alkoxy carbonyl-substituted lower alkyl group;

A is a lower alkylene group; and  
n is 0 or 1.

Claim 2. (currently amended) The method according to Claim 1, wherein  $R^3$  is an indolyl group, said indolyl group may have 1 to 3 substituents selected from the group consisting of: a group of the formula  $-B-R^4$ , (wherein B is a lower alkylene group;  $R^4$  is a 5- to 11-membered saturated or unsaturated heterocyclic group of single ring or binary ring, having 1 to 4 hetero atoms selected from the group consisting of a nitrogen atom, oxygen atom and sulfur atom, (said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a halogen atom, a lower alkyl group, a lower alkoxy group and oxo group) or a group of the formula  $-NR^5R^6$  (wherein  $R^5$  and  $R^6$  are each the same or different, and a hydrogen atom, a lower alkyl group, a cycloalkyl group, a pyridylcarbonyl group, an isoxazolylcarbonyl group which may have 1 to 3 lower alkyl groups as the substituents, a pyrrolylcarbonyl group or an amino-substituted lower alkyl group which may have a lower alkyl group as the substituent; further  $R^5$  and  $R^6$  may form 5- to 6membered saturated heterocyclic group by combining to each other, together with the adjacent nitrogen atom being bonded thereto, further with or without other nitrogen atom or oxygen atom; said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a hydroxy group and a phenyl group) [group)); a lower alkenyl group; a lower alkoxycarbonyl group; a phenoxy-lower alkyl group which may have cyano group as the substituents; a halogen-substituted lower alkyl group; and a lower alkoxycarbonyl-substituted lower alkyl group.

Claim 3. (currently amended) A method for inhibiting the growth of neoplastic cells comprising exposing the cells to a growth inhibiting effective amount of a compound of Formula I or pharmaceutically acceptable salt thereof:



(I)

wherein [R<sub>1</sub> is] R<sup>1</sup> is a hydrogen atom or a halogen atom;

[R<sub>2</sub> is] R<sup>2</sup> is a phenyl-lower alkyl group;

[R<sub>3</sub> is] R<sup>3</sup> is a heterocyclic group selected from the group consisting of an indolyl group, indolinyl group, 1H-indazolyl group, 2(1H)-quinolinonyl group, 3,4-dihydro-2(1H)-quinolinonyl group and 3,4-dihydro-1,4(2H)-benzoxazinyl group, said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a group of the formula -B-R<sup>4</sup>, (B is a lower alkylene group; R<sup>4</sup> is a 5-to 11-membered saturated or unsaturated heterocyclic group of single ring or binary ring, having 1 to 4 hetero atoms selected from the group consisting of a nitrogen atom, oxygen atom and sulfur atom, (said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a halogen atom, a lower alkyl group, a lower alkoxy group and oxo group) or a group of the formula -NR<sup>5</sup>R<sup>6</sup> (R<sup>5</sup> and R<sup>6</sup> are each the same or different, and a hydrogen atom, a lower alkyl group, a cycloalkyl group, a pyridyl-carbonyl group, an isoxazolylcarbonyl group which may have 1 to 3 lower alkyl groups as the substituents, a pyrrolylcarbonyl group, or an amino-substituted lower alkyl group which may have a lower alkyl group as the substituent; further R<sup>5</sup> and R<sup>6</sup> may form 5- to 6-membered saturated heterocyclic group by combining to each other, together with the adjacent nitrogen atom being bonded thereto, further with or without other nitrogen atom or oxygen atom; said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a hydroxy group and a phenyl group)); a lower alkenyl group; a lower alkoxy carbonyl group; a phenoxy-lower alkyl group which may have cyano group as the substituents; a halogen-substituted lower alkyl group; and a lower alkoxy carbonyl-substituted lower alkyl group;

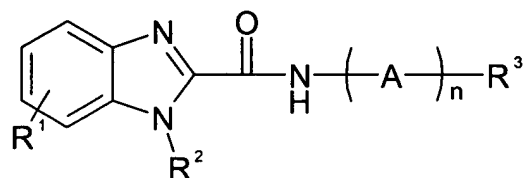
A is a lower alkylene group; and

n is 0 or 1.

Claim 4. (currently amended) The method according to Claim 3, wherein R<sup>3</sup> is an indolyl group, said indolyl group may have 1 to 3 substituents selected from the group consisting of: a group of the formula -B-R<sup>1</sup>, (B is a lower alkylene group; R<sup>4</sup> is a 5- to 11-membered saturated or unsaturated heterocyclic group of single ring or binary ring, having 1 to 4 hetero atoms selected from the group consisting of a nitrogen atom, oxygen

atom and sulfur atom, (said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a halogen atom, a lower alkyl group, a lower alkoxy group and oxo group) or a group of the formula  $-NR^5R^6$  (wherein  $R^5$  and  $R^6$  are each the same or different, and a hydrogen atom, a lower alkyl group, a cycloalkyl group, a pyridyl-carbonyl group, an isoxazolylcarbonyl group which may have 1 to 3 lower alkyl groups as the substituents, a pyrrolylcarbonyl group or an amino-substituted lower alkyl group which may have a lower alkyl group as the substituent; further  $R^5$  and  $R^6$  may form 5- to 6membered saturated heterocyclic group by combining to each other, together with the adjacent nitrogen atom being bonded thereto, further with or without other nitrogen atom or oxygen atom; said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a hydroxy group and a phenyl [group])) group); a lower alkenyl group; a lower alkoxy carbonyl group; a phenoxy-lower alkyl group which may have cyano group as the substituents; a halogen-substituted lower alkyl group; and a lower alkoxy carbonyl-substituted lower alkyl group.

Claim 5. (currently amended) A method for regulating apoptosis in human cells comprising exposing said cells to an effective amount of a compound of the formula:



wherein  $[R_1]$  is  $R^1$  is a hydrogen atom or a halogen atom;

$[R_2]$  is  $R^2$  is a phenyl-lower alkyl group;

$[R_3]$  is  $R^3$  is a heterocyclic group selected from the group consisting of an indolyl group, indolyl group, 1H-indazolyl group, 2(1H)-quinolinonyl group, 3,4-dihydro-2(1H)-quinolinonyl group and 3,4-dihydro- 1,4(2H)-benzoxazinyl group, said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a group of the formula  $-B-R^4$ , ([.a] wherein B is a lower alkylene group;  $R^4$  is a 5-to 11-membered saturated or unsaturated heterocyclic group of single ring or binary ring, having 1 to 4 hetero atoms selected from the group consisting of a nitrogen atom, oxygen atom and sulfur atom, (said heterocyclic group may have 1 to 3 substituents selected from

the group consisting of a halogen atom, a lower alkyl group, a lower alkoxy group and oxo group) or a group of the formula  $-NR^5R^6$  (wherein  $R^5$  and  $R^6$  are each the same or different, and each is a hydrogen atom, a lower alkyl group, a cycloalkyl group, a pyridyl-carbonyl group, an isoxazolylcarbonyl group which may have 1 to 3 lower alkyl groups as the substituents, a pyrrolylcarbonyl group, or an amino-substituted lower alkyl group which may have a lower alkyl group as the substituent; further  $R^5$  and  $R^6$  may form 5- to 6-membered saturated heterocyclic group by combining to each other, together with the adjacent nitrogen atom being bonded thereto, further with or without other nitrogen atom or oxygen atom; said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a hydroxy group and a phenyl [group))) group); a lower alkenyl group; a lower alkoxycarbonyl group; a phenoxy-lower alkyl group which may have cyano group as the substituents; a halogen-substituted lower alkyl group; and a lower alkoxycarbonyl-substituted lower alkyl group;

A is a lower alkylene group; and

n is 0 or 1.

Claim 6. (currently amended) The method according to Claim 5, wherein  $R^3$  is an indolyl group, said indolyl group may have 1 to 3 substituents selected from the group consisting of: a group of the formula  $-B-R^4$ , (wherein B is a lower alkylene group;  $R^4$  is a 5- to 11-membered saturated or unsaturated heterocyclic group of single ring or binary ring, having 1 to 4 hetero atoms selected from the group consisting of a nitrogen atom, oxygen atom and sulfur atom, (said heterocyclic group may have 1 to 3 substituents selected from the group consisting of a halogen atom, a lower alkyl group, a lower alkoxy group and oxo group) or a group of the formula  $-NR^5R^6$  (wherein  $R^5$  and  $R^6$  are each the same or different, and each is a hydrogen atom, a lower alkyl group, a cycloalkyl group, a pyridylcarbonyl group, an isoxazolylcarbonyl group which may have 1 to 3 lower alkyl groups as the substituents, a pyrrolylcarbonyl group or an amino-substituted lower alkyl group which may have a lower alkyl group as the substituent; further  $R^5$  and  $R^6$  may form 5- to 6membered saturated heterocyclic group by combining to each other, together with the adjacent nitrogen atom being bonded thereto, further with or without other nitrogen atom or oxygen atom; said heterocyclic group may have 1 to 3 substituents selected from

the group consisting of a hydroxy group and a phenyl [group))] group); a lower alkenyl group; a lower alkoxycarbonyl group; a phenoxy-lower alkyl group which may have cyano group as the substituents; a halogen-substituted lower alkyl group; and a lower alkoxycarbonyl-substituted lower alkyl group.